## AMENDMENTS TO THE CLAIMS:

This listing of the claims below will replace all prior versions and listing of claims:

## Listing of Claims

Claims 1-43 (Cancelled).

- 44. (Currently amended) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:
  - (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene encoding chymosin from a bovine or Camelidae-species or Camelus dromedarius.
  - (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.8 by addition of <del>lactic acid, acetic acid, propionic acid, or citric acid, and</del>
  - (iii) subjecting said medium to a pH in the range of 1.0 to 1.8 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.
- 45. (Previously presented) The method according to claim 44, wherein at least 90% of said glucoamylase activity is inactivated.
- 46. (Previously presented) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is a medium derived from the cultivation of an organism that during its cultivation produces said chymosin activity and said glucoamylase activity.
- 47. (Previously presented) The method according to claim 44, wherein the bacterial species is a gram negative bacterial species or a gram positive species.
- 48. (Previously presented) The method according to claim 47, wherein the bacterial species is E. coli or Bacillus.

- 49. (Previously presented) The method according to claim 44, where the yeast species is Saccharomyces cerevisiae, a methylotrophic yeast species or a Klyweromyces species.
- 50. (Previously presented) The method according to claim 44, wherein the species of filamentous fungi is an Aspergillus species, a Cryphonectria species, a Fusarium species, a Rhizomucor species or a Trichoderma species.
- (Previously presented) The method of claim 50, wherein said Aspergillus species is Aspergillus niger var. awamori.
- 52. (Previously presented) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH in the range of 1.5 to 1.8.
- 53. (Previously presented) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH between 1.5 to 1.7.
- 54. (Previously presented) The method according to claim 44, wherein the medium having a pH of 2.0 or higher is subjected to a pH of approximately 1.8.
- 55. (Previously presented) The method according to claim 44, wherein said period of time is in the range of 0.1 minutes to 48 hours.
- 56. (Previously presented) The method according to claim 44, wherein lowering of the pH in step (ii) is performed by addition of acetic acid.
- (Previously presented) The method of claim 44, wherein the gene encoding chymosin is from Camelus dromedarius.
- 58. (Previously presented) The method of claim 44, wherein at least 85% of the chymosin activity is maintained in step (iii).
- (Previously presented) The method of claim 44, wherein the gene encoding chymosin is from a bovine species.
- 60. (Currently amended) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:

- (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene encoding chymosin from a bovine or Camelidae species or Camelus dromedarius.
- (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.7 by addition of an inorganic acid, and
- (iii) subjecting said medium to a pH in the range of 1.0 to 1.7 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.
- (Previously presented) The method according to claim 60, wherein at least 90% of said glucoamylase activity is inactivated.
- 62. (Previously presented) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is a medium derived from the cultivation of an organism that during its cultivation produces said chymosin activity and said glucoamylase activity.
- 63. (Previously presented) The method according to claim 60, wherein the bacterial species is a gram negative bacterial species or a gram positive species.
- 64. (Previously presented) The method according to claim 63, wherein the bacterial species is E. coli or Bacillus.
- (Previously presented) The method according to claim 60, where the yeast species is Saccharomyces cerevisiae, a methylotrophic yeast species or a Klyuveromyces species.
- 66. (Previously presented) The method according to claim 60, wherein the species of filamentous fungi is an Aspergillus species, a Cryphonectria species, a Fusarium species, a Rhizomucor species or a Trichoderma species.
- 67. (Previously presented) The method of claim 66, wherein said Aspergillus species is

Aspergillus niger var. awamori.

- 68. (Previously presented) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH in the range of 1.5 to 1.7.
- (Previously presented) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH between 1.5 to 1.7.
- (Previously presented) The method according to claim 60, wherein the medium having a pH of 2.0 or higher is subjected to a pH of approximately 1.7.
- (Previously presented) The method according to claim 60, wherein said period of time is in the range of 0.1 minutes to 48 hours.
- (Previously presented) The method according to claim 60, wherein the yeast species is selected from Pichia pastoris and Klyuveromyces lactis.
- (Previously presented) The method of claim 60, wherein the gene encoding chymosin is from Camelus dromedarius.
- (Previously presented) The method of claim 60, wherein at least 85% of the chymosin activity is maintained in step (iii).
- 75. (Previously presented) The method of claim 60, wherein the gene encoding chymosin is from a bovine species.
- (Previously presented) The method of claim 60, wherein the inorganic acid is hydrochloric acid, phosphoric acid, or sulfuric acid.
- (Currently amended) The method of claim 60, wherein the glucoamylase is derived from culture of an Aspergillus species.
- 78. (Currently amended) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:
  - (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity

- and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene encoding chymosin from a bovine or Camelidae species or Camelia dromedarius,
- (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.8 by addition of inorganic acid and an organic acid, wherein the organic acid is acetic acid or propionic acid; and
- (iii) subjecting said medium to a pH in the range of 1.0 to 1.8 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.
- 79. (New) A method for reducing the glucoamylase activity in a milk clotting composition comprising the steps of:
  - (i) providing a medium having a pH of 2.0 or higher that comprises chymosin activity and glucoamylase activity, wherein the medium having a pH of 2.0 or higher is derived from the cultivation of an organism that is a bacterial species, a yeast species, or a species of filamentous fungi, wherein the organism comprises a gene encoding chymosin from a bovine species or Camelus dromedarius,
  - (ii) lowering the pH of said medium having a pH of 2.0 or higher to between 1.0 and 1.8 by addition of lactic acid, and
  - (iii) subjecting said medium to a pH in the range of 1.0 to 1.7 for a period of time sufficient to inactivate at least 50% of said glucoamylase activity while maintaining at least 75% of said chymosin activity.